

## Dielectric properties of 1:1 ternary $\text{Pb}_2\text{B}^{3+}\text{B}^{5+}\text{O}_6$ perovskite ceramics sintered from mechanochemically synthesized nanopowders

S.I. Raevskaya<sup>1</sup>, A.A. Gusev<sup>2</sup>, V.P. Isupov<sup>2</sup>, I.P. Raevski<sup>1</sup>, V.V. Titov<sup>1</sup>, M.A. Malitskaya<sup>1</sup>,  
G.R. Li<sup>3</sup>, S.P. Kubrin<sup>1</sup>, E.I. Sitalo<sup>1</sup>

<sup>1</sup>Research Institute of Physics and Faculty of Physics, Southern Federal University, 344090, Rostov-on-Don, Russia  
e-mail: sveta.raevskaya@mail.ru

<sup>2</sup>Institute of Solid State Chemistry and Mechanochemistry SB RAS, 630128, Novosibirsk, Russia

<sup>3</sup>Shanghai Institute of Ceramics, Chinese Academy of Sciences, 200050, Shanghai, China

Ferroelectric 1:1 ternary perovskite oxides of the  $\text{Pb}_2\text{B}^{3+}\text{B}^{5+}\text{O}_6$  type, are widely used as a basis or components of various functional materials exhibiting giant dielectric, electrostrictive, pyroelectric, and piezoelectric responses. Properties of these oxides depend crucially on the ordering degree of  $\text{B}^{3+}$  and  $\text{B}^{5+}$  cations. While in the highly-ordered state a sharp ferroelectric or antiferroelectric phase transition takes place, in the disordered state a relaxor state is observed, characterized by a diffuse and frequency-dependent maximum of the dielectric permittivity.

Recently we have found out that the use of mechanochemical synthesis enables one to modify substantially relaxor properties of ceramics of the disordered perovskites  $\text{Pb}_2\text{Fe}^{3+}\text{Ta}^{5+}\text{O}_6$  and  $\text{Pb}_2\text{MgNb}_2\text{O}_9$  [1,2]. This effect is believed to be caused by the changes of the degree of the short-range ordering. Such explanation is supported by changes in the magnetic phase transition temperature of  $\text{Pb}_2\text{Fe}^{3+}\text{B}^{5+}\text{O}_6$  powders ( $\text{B}^{5+}$  - Nb, Ta) obtained by mechanochemical synthesis [3]. In the present study we investigated the effect of the conditions of the mechanochemical synthesis on the structure and dielectric properties of several  $\text{Pb}_2\text{B}^{3+}\text{B}^{5+}\text{O}_6$  ceramics ( $\text{B}^{3+}$  - Sc, In, Yb and  $\text{B}^{5+}$  - Nb, Ta). These ternary perovskite oxides of the 1:1 type are known to possess a high degree of long-range ordering of  $\text{B}^{3+}$  and  $\text{B}^{5+}$  ions when synthesized by a usual solid state route. Mechanochemical synthesis was carried out using high-energy planetary-centrifugal mill AGO-2 with steel balls and jars under a ball acceleration of 40g.

It was found out that high-energy mechanical activation during mechanochemical synthesis stimulates disordering of  $\text{B}^{3+}$  and  $\text{B}^{5+}$  cations in ceramics of the  $\text{Pb}_2\text{B}^{3+}\text{B}^{5+}\text{O}_6$  perovskites studied. Using mechanochemical synthesis disordered modifications of  $\text{Pb}_2\text{B}^{3+}\text{B}^{5+}\text{O}_6$  ( $\text{B}^{3+}$  - In, Yb and  $\text{B}^{5+}$  - Nb, Ta) were obtained for the first time without the use of any additives. In some cases it appeared to be possible to vary within a wide range compositional ordering degree of the  $\text{B}^{3+}$  and  $\text{B}^{5+}$  cations and correspondingly both the temperature and diffusion of dielectric permittivity maxima in ceramics by changing the regimes of the mechanochemical synthesis and/or using different starting materials (either oxides or preliminary synthesized  $\text{B}^{3+}\text{B}^{5+}\text{O}_4$  precursors). The parameters of the dielectric spectra of the samples exhibiting relaxor-like behaviour also depend on the conditions of the mechanochemical synthesis

This study was supported by RFBR (project 17-03-01293\_a) and by the Ministry of Education and Science of the Russian Federation (project 3.1649.2017/4.6).

1. A.A. Gusev, S.I. Raevskaya, V.V. Titov, et al., *Ferroelectrics* **475**, 41 (2015).
2. S.I. Raevskaya, A.A. Gusev, V.P. Isupov, et al., *Ferroelectrics* **525**, 37 (2018).
3. A.A. Gusev, S.I. Raevskaya, V.V. Titov, et al., *Ferroelectrics* **496**, 231 (2016).